## IN THE CLAIMS:

1. A compound comprising:

$$\begin{split} &H_xCs_{(4-x)}Ge_4S_{10}\dot{\cdot}yH_2S\dot{\cdot}zH_2O,\, H_xRb_{(4-x)}Ge_4S_{10}\dot{\cdot}zH_2O \text{ or } H_xK_{(4-x)}Ge_4S_{10}\dot{\cdot}zH_2O,\\ &_{x)}Ge_4S_{10}\dot{\cdot}zH_2O, \end{split}$$

wherein

 $0 < x \le 4;$  $0 \le y \le 1;$  and

 $z \ge 0$ .

- 2. The compound of claim 1 wherein x = 4.
- 3. The compound of claim 1 wherein z = 0.
- 4. The compound of claim 3 comprising a compound having a double-decker structural unit
- 5. The compound of claim 4 wherein the double-decker compound is  $H_2Ge_4S_9$ .
- 6. The compound of claim 3 comprising a compound having an adamantane-like structural unit.
- 7. The compound of claim 6 wherein the adamantane-like compound is  $H_4Ge_4S_{10}$ .
- 8. The compound of claim 1 wherein z > 0.

- 9. The compound of claim 8 comprising a compound having a double-decker structural unit
- 10. The compound of claim 9 wherein the double-decker compound is  $H_4Ge_4S_{10}$  x $H_2O$  wherein 0 < x < 9.
- 11. The compound of claim 8 comprising a compound having an adamantane-like structural unit.
- 12. The compound of claim 11 wherein the adamante compound is  $H_4Ge_4S_{10}$  x $H_2O$  wherein 0 < x < 9.
- 13. The compound of claim 1 wherein the proton conductivity is between about  $10^{-8}$  S/cm and  $10^{-1}$  S/cm within a temperature range of between about -50 and 500 °C.
- 14. A compound comprising:

 $H_xMoS_{2+x/2+z}$ ,

wherein

 $1 \le x \le 2$ ; and

 $0.5 \le z \le 1$ .

- 15. The compound of claim 14 comprising a thiomolybdic acid.
- 16. The compound of claim 15 wherein the thiomolybdic acid is anhydrous or hydrated.
- 17. The compound of claim 14 wherein the thiomolybdic acid is an anhydrous thiomolybdic acid having a conductivity value of 10<sup>-8</sup> S/cm and 10<sup>-1</sup> S/cm within a temperature range of between about -50 and 500 °C.

## 18. A method comprising:

reacting liquid H<sub>2</sub>S, H<sub>2</sub>Se or H<sub>2</sub>Te with a precursor selected from the group consisting of chalcogenides, halides, and any combination thereof, to produce a protonated chalcogenide or unprotonated crystalline sulfide, selenide or telluride..

- 19. The method of claim 18 wherein the precursor is selected from the group consisting of an oxide, sulfide, hydroxide, fluoride, chloride, iodide, bromide, and any combination thereof, to produce a thio-acid, a hydrosulfide or a sulfide.
- 20. The method of claim 18 wherein the reacting step occurs in a closed oneend tube located inside a reaction vessel.
- 21. The method of claim 20 wherein the reaction vessel has a reusable sealing mechanism.
- 22. The method of claim 20 further comprising:

  placing the precursor in the reaction vessel;

  drawing a vacuum inside the reaction vessel;

  cooling the reaction vessel;

  filling the reactor with the H<sub>2</sub>S, H<sub>2</sub>Se or H<sub>2</sub>Te gas;

  allowing the H<sub>2</sub>S, H<sub>2</sub>Se or H<sub>2</sub>Te gas to liquefy and/or solidify; and

  allowing the reaction to proceed wherein the protonated chalcogenide or

  unprotonated sulfide, selenide or telluride is produced.
- 23. The method of claim 22 wherein the reaction vessel is cooled to below about -60 °C.
- 24. The method of claim 22 further comprising opening the reactor inside an oxygen and water free glove box after the reaction is complete.

- 25. The method of claim 24 further comprising thermally mixing the protonated chalcogenide with other chalcogenide materials in an evacuated silica tube to produce a glass, ceramic or a glass/ceramic membrane material.
- 26. The method of claim 25 wherein the protonated chalcogenide is a thio-acid or hydrosulfide which is mixed with other sulfide materials.
- 27. The method of claim 24 wherein a glass material is produced by combining two or more of the chalcogenides.
- 28. The method of claim 24 wherein a ceramic material is produced by combining two or more of the chalcogenides.
- 29. The method of claim 24 wherein a glass/ceramic material is produced by combining two or more of the chalcogenides.
- 30. The method of claim 24 further comprising reacting the glass, ceramic or glass/ceramic material with liquid H<sub>2</sub>S, H<sub>2</sub>Se or H<sub>2</sub>Te to produce a chemically and thermally stable membrane material.
- 31. A method for producing a chemically and thermally stable membrane material comprising:

providing a glass phase, ceramic phase or glass/ceramic phase material; and

allowing the glass phase, ceramic phase or glass/ceramic phase material to protonate in liquid H<sub>2</sub>S, H<sub>2</sub>Se or H<sub>2</sub>Te to produce a chemically and thermally stable membrane material.

- 32. The method of claim 31 wherein the glass phase, ceramic phase or glass/ceramic phase material is a chalcogenide.
- 33. The method of claim 32 wherein the chalcogenide is selected from the group consisting of oxide, sulfide and oxysulfide.
- 34. A chemically and thermally stable membrane comprising a glass material, ceramic material or glass/ceramic material adaptable for use in a fuel cell at a temperature of between about -60 °C and 700 °C, the material having a proton conductivity of between about 10<sup>-6</sup> to 10<sup>-1</sup> S/cm.
- 35. The membrane of claim 34 wherein the temperature is greater than 100 °C and less than about 500 °C.